In claim 8, at line 1, after "inhibitor" please insert --composition--.

In claim 9, at line 1, after "inhibitor" please insert -- composition--.

In claim 10, at line 1, after "inhibitor" please insert --composition--.

In claim 11, at line 1, after "inhibitor" please insert -- composition--.

In claim 12, at line 1, after "inhibitor" please insert -- composition--.

In claim 14, at line 1, after "inhibitor" please insert --composition--.

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In claim 16, at line 1, please delete "inhibitor of" and after "production" please insert --inhibitor composition--.

In claim 17, at line 1, please delete "inhibitor of" and after "production" please insert --inhibitor composition--.

In claim 18, at line 1, please delete "inhibitor of" and after "production" please insert --inhibitor composition--.

In claim 19, at line 1, please delete "inhibitor of" and after "production" please insert --inhibitor composition--.

In claim 20, at line 1, please delete "inhibitor of" and after "production" please insert --inhibitor composition--.

In claim 21, at line 1, please delete "inhibitor of" and after "production" please insert --inhibitor composition--.

In claim 22, at line 1, please delete "inhibitor of" and after "production" please insert --inhibitor composition--.

In claim 23, at line 1, please delete "inhibitor of" and after "production" please insert --inhibitor composition--.

In claim 24, at line 1, please delete "inhibitor of" and after "production" please insert --inhibitor composition--.

In claim 25, at line 1, please delete "inhibitor of" and after "production" please insert --inhibitor composition--.

In claim 26, at line 1, please delete "inhibitor of" and after "production" please insert --inhibitor composition--.

In claim 27, at line 1, please delete "inhibitor of" and after "production" please insert --inhibitor composition--.

In claim 28, at line 1, please delete "inhibitor of" and after "production" please insert --inhibitor composition--.

Please add the following new claims 38-41:

--38. A method for inhibiting NF-κB comprising administering to a patient in need of NF-κB inhibition a benzoquinone derivative represented by the following general formula (1):

$$\begin{array}{c}
O \\
R_3 \\
CH_2-Z-(CH_2)_n-R_4
\end{array}$$

wherein

 R_1 , R_2 and R_3 are each independently a hydrogen atom, an alkyl group having 1 to 5 carbons, or an alkoxy group having 1 to 5 carbons;

R₄ is a hydrogen atom, a hydroxymethyl group, an alkyl group, or a carboxyl group which is optionally esterified or amidated;

and, n is

an integer from 0 to 6, or its hydroquinone form, or a pharmaceutically acceptable salt thereof.

39. A method for preventing or treating diseases caused by the activation of NFκB comprising administering to a patient a benzoquinone derivative represented by the following general formula (1): C5 cont B'

$$\begin{array}{c} O \\ R_3 \\ CH_2-Z-(CH_2)_n-R_4 \end{array}$$

wherein

 R_1 , R_2 , and R_3 are each independently a hydrogen atom, an alkyl group having 1 to 5 carbons, or an alkoxy group having 1 to 5 carbons;

R₄ is a hydrogen atom, a hydroxymethyl group, an alkyl group, or a carboxyl group which is optionally esterified or amidated;

Z is

and, n is an integer from 0 to 6,

or its hydroquinone form, or a pharmaceutically acceptable salt thereof.

40. A method for inhibiting TNF- α production comprising administering to a patient in need of TNF- α inhibition a benzoquinone derivative represented by the following general formula (1):

$$R_{3}$$
 $CH_{2}-Z-(CH_{2})_{n}-R_{4}$

wherein R_1 , R_2 and R_3 are each independently a hydrogen atom, an alkyl group having 1 to 5 carbons, or an alkoxy group having 1 to 5 carbons;

R₄ is a hydrogen atom, a hydroxymethyl group, an alkyl group, or a carboxyl group which is optionally esterified or amidated;

Z is

a c

and, n is an integer from 0 to 6,

or its hydroquinone form, or a pharmaceutically acceptable salt thereof.

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41. A method for preventing or treating diseases caused by the excessive production of TNF-α comprising administering to a patient a benzoquinone derivative represented by the following general formula (1):

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$$R_{3}$$
 $CH_{2}-Z-(CH_{2})_{n}-R_{4}$

wherein R_1 , R_2 and R_3 are each independently a hydrogen atom, an alkyl group having 1 to 5 carbons, or an alkoxy group having 1 to 5 carbons;

R₄ is a hydrogen atom, a hydroxymethyl group, an alkyl group, or a carboxyl group which is optionally esterified or amidated;

Z is